BIMECTIN PLUS 10/100 MG/ML SOLUTION FOR INJECTION FOR CATTLE

Authorised

- Ivermectin
- Clorsulon

Product identification

Medicine name:
BIMECTIN PLUS 10/100 MG/ML SOLUTION FOR INJECTION FOR CATTLE
Bimectin plus 10/100 mg/ml solução injetável para bovinos

Active substance:
Ivermectin
Clorsulon

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Ivermectin

10.00 milligram(s) / 1.00 millilitre(s)

Clorsulon
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Cattle**
  - Meat and offal. 66 day
  - Milk. no withdrawal period

Milk: Do not use in cattle producing milk for human consumption. Do not use in non-lactating dairy cows including pregnant heifers within 60 days of calving.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AA51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Portugal

Package description:
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
21/01/2011

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
307/01/11RFVPT

Date of authorisation status change:
27/05/2025

Reference member state:
France

Procedure number:
FR/V/0337/001

Concerned member states:
Belgium Denmark Germany Italy Poland Portugal Romania Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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